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FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO	
Dan W. Denney JR.	GENITOPE-06493	5113	
1	EXAMINER		
MEDLEN & CARROLL, LLP			
	ART UNIT	PAPER NUMBER	
	1642	-	
),,		Dan W. Denney JR. GENITOPE-06493 EXAM YAEN, CHRI	

DATE MAILED: 08/24/2004

Please find below and/or attached an Office communication concerning this application or proceeding.

 		Applicati	ion No.	Applicant(s)			
		09/925,1	92	DENNEY, DAN W.			
Office Action Summary		Examine		Art Unit			
		Christoph	ner H Yaen	1642			
	The MAILING DATE of this commu			th the correspondence add	ress		
Period fo	• •						
THE - External control	ORTENED STATUTORY PERIOD MAILING DATE OF THIS COMMUN ensions of time may be available under the provision of SIX (6) MONTHS from the mailing date of this come period for reply specified above is less than thirty. Despriod for reply is specified above, the maximum is ure to reply within the set or extended period for repreply received by the Office later than three months and patent term adjustment. See 37 CFR 1.704(b).	NICATION. as of 37 CFR 1.136(a). In no expression. (30) days, a reply within the statatutory period will apply and ly will, by statute, cause the ap	vent, however, may a ratutory minimum of third will expire SIX (6) MON plication to become AE	eply be timely filed by (30) days will be considered timely. ITHS from the mailing date of this com SANDONED (35 U.S.C. § 133).	nmunication.		
Status							
1)⊠	Responsive to communication(s) fi	led on 09 August 200	1.				
•	This action is FINAL .	2b) This action is i					
3)□	Since this application is in condition	n for allowance excep	t for formal matt	ers, prosecution as to the r	merits is		
	closed in accordance with the practice under Ex parte Quayle, 1935 C.D. 11, 453 O.G. 213.						
Disposit	ion of Claims						
·		application					
7)23	4) Claim(s) 1-24 is/are pending in the application. 4a) Of the above claim(s) is/are withdrawn from consideration.						
5)[]	Claim(s) is/are allowed.						
	_						
	Claim(s) <u>1-24</u> are subject to restrict	tion and/or election re	quirement.				
Applicat	ion Papers						
	•	he Evaminer					
9) The specification is objected to by the Examiner. 10) The drawing(s) filed on is/are: a) accepted or b) objected to by the Examiner.							
10)[Applicant may not request that any obj						
	Replacement drawing sheet(s) including				R 1.121(d).		
11)	The oath or declaration is objected						
,—	under 35 U.S.C. § 119	•					
•	Acknowledgment is made of a claim	a for foreign priority ur	nder 35 II S.C. 8	: 119/a) ₋ (d) or (f)			
		ir for foreign priority ar	ide: 55 0.5.0. §	7 113(a)-(d) or (i).			
a)	a) All b) Some * c) None of: 1. Certified copies of the priority documents have been received.						
	2. Certified copies of the priority			nolication No			
	3. Copies of the certified copies				stage		
	application from the Internati	, ,		TOURIST THE TRANSPORT	90		
* (See the attached detailed Office acti	•		received.			
			•				
Attach	54(a)						
Attachmer 1) Notice	ce of References Cited (PTO-892)		4) Interview 9	Summary (PTO-413)			
	ce of Draftsperson's Patent Drawing Review	(PTO-948)	Paper No(s	s)/Mail Date			
3) Infor	mation Disclosure Statement(s) (PTO-1449 or er No(s)/Mail Date		5) Notice of I	nformal Patent Application (PTO- 	152)		

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DETAILED ACTION

Election/Restrictions

- 1. Restriction to one of the following inventions is required under 35 U.S.C. 121:
 - I. Claims 1-6, drawn to a multivalent vaccine comprising at least two recombinant variable regions, classified in class 530, subclass 387.1.
 - II. Claims 7-20, drawn to a method of producing a vaccine for the treatment of B-cell lymphoma, classified in class 435, subclass 69.1. *If applicant elects this group for prosecution on the merits, applicant is required to select ONE inhibitable enzyme from dihydrofolate reductase, glutamine synthetase, adenosine deaminase, or asparagines synthetase. This selection should <u>not</u> be construed as an election of species (see paragraph 7 below).*
 - III. Claims 21-24, drawn to a method of treating B-cell lymphoma comprising the administration of a multivalent vaccine of group I, classified in class 424, subclass 178.1.

The inventions are distinct, each from the other because of the following reasons:

2. Inventions II and III are unrelated. Inventions are unrelated if it can be shown that they are not disclosed as capable of use together and they have different modes of operation, different functions, or different effects (MPEP § 806.04, MPEP § 808.01). In the instant case the different inventions differ one from the other in that the methods are

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used for either production or administration of a multivalent composition. Thus the method steps and purpose of use are different and distinct.

- 3. Inventions II and I are related as process of making and product made. The inventions are distinct if either or both of the following can be shown: (1) that the process as claimed can be used to make other and materially different product or (2) that the product as claimed can be made by another and materially different process (MPEP § 806.05(f)). In the instant case the product can be made through other means such as through the extraction and isolation of phage antibodies.
- 4. Inventions I and III are related as product and process of use. The inventions can be shown to be distinct if either or both of the following can be shown: (1) the process for using the product as claimed can be practiced with another materially different product or (2) the product as claimed can be used in a materially different process of using that product (MPEP § 806.05(h)). In the instant case the method can be accomplished by administering peptides or other organic molecules that are capable of inhibiting the growth of malignant B-cells.
- 5. Because these inventions are distinct for the reasons given above and have acquired a separate status in the art as shown by their different classification, restriction for examination purposes as indicated is proper.
- 6. This application contains claims directed to the following patentably distinct species of the claimed invention:

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- a. If applicant elects group I, applicant must elect a <u>single</u> immune enhancing cytokine from claim 4 (i.e. granulocyte-macrophage colony stimulating factor, IL-2, or IL-4)
- b. If applicant elects group II, applicant must elect a <u>single</u> inhibitor from claim 20 (i.e. methotrexate, 2'-deoxycoformycin, methionine sulphoximine, albizziin, or beta-aspartyl hydroxamate).

Applicant is required under 35 U.S.C. 121 to elect a single disclosed species for prosecution on the merits to which the claims shall be restricted if no generic claim is finally held to be allowable. Currently, claims 3 and 19 are generic.

Applicant is advised that a reply to this requirement must include an identification of the species that is elected consonant with this requirement, and a listing of all claims readable thereon, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered nonresponsive unless accompanied by an election.

Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which are written in dependent form or otherwise include all the limitations of an allowed generic claim as provided by 37 CFR 1.141. If claims are added after the election, applicant must indicate which are readable upon the elected species. MPEP § 809.02(a).

Should applicant traverse on the ground that the species are not patentably distinct, applicant should submit evidence or identify such evidence now of record showing the species to be obvious variants or clearly admit on the record that this is the

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case. In either instance, if the examiner finds one of the inventions unpatentable over the prior art, the evidence or admission may be used in a rejection under 35 U.S.C. 103(a) of the other invention.

Applicant is advised that the reply to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed (37 CFR 1.143).

7. Upon election of Group II, Applicants are additionally required to elect a single inhibitable enzyme, as indicated above as they apply to group(s). The recited enzymes all have different structures one from other and the search for different enzymes as they apply to the method would be unduly burdensome. This requirement is not to be construed as a requirement for an election of species, since each of enzymes recited in alternative form is not a member of a single genus of invention, but constitutes an independent and patentably distinct invention.

Note:

The examiner has required restriction between product and process claims. Where applicant elects claims directed to the product, and a product claim is subsequently found allowable, withdrawn process claims that depend from or otherwise include all the limitations of the allowable product claim will be rejoined in accordance with the provisions of MPEP § 821.04. Process claims that depend from or otherwise include all the limitations of the patentable product will be entered as a matter of right if the amendment is presented prior to final rejection or allowance, whichever is earlier. Amendments submitted after final

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rejection are governed by 37 CFR 1.116; amendments submitted after allowance are governed by 37 CFR 1.312.

In the event of rejoinder, the requirement for restriction between the product claims and the rejoined process claims will be withdrawn, and the rejoined process claims will be fully examined for patentability in accordance with 37 CFR 1.104. Thus, to be allowable, the rejoined claims must meet all criteria for patentability including the requirements of 35 U.S.C. 101, 102, 103, and 112. Until an elected product claim is found allowable, an otherwise proper restriction requirement between product claims and process claims may be maintained. Withdrawn process claims that are not commensurate in scope with an allowed product claim will not be rejoined. See "Guidance on Treatment of Product and Process Claims in light of *In re Ochiai, In re Brouwer* and 35 U.S.C. § 103(b)," 1184 O.G. 86 (March 26, 1996). Additionally, in order to retain the right to rejoinder in accordance with the above policy, Applicant is advised that the process claims should be amended during prosecution either to maintain dependency on the product claims or to otherwise include the limitations of the product claims. **Failure to do so may result in a loss of the right to rejoinder.**

Further, note that the prohibition against double patenting rejections of 35 U.S.C. 121 does not apply where the restriction requirement is withdrawn by the examiner before the patent issues. See MPEP § 804.01.

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Any inquiry concerning this communication or earlier communications from the examiner should be directed to Christopher H Yaen whose telephone number is 571-272-0838. The examiner can normally be reached on Monday-Friday 9-5.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Jeffrey Siew can be reached on 571-272-0787. The fax phone number for the organization where this application or proceeding is assigned is 703-872-9306.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

Christopher Yaen Art Unit 1642 August 9, 2004

GARY NICKOL
PRIMARY EXAMINER

Mary & Mikel